



RECEIVED

OCT 03 2002

16141

## TECH CENTER 1600/2900

Approved for use through 10/31/2002. OMB 0651-0032

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

# FEE TRANSMITTAL for FY 2002

Patent fees are subject to annual revision.

 Applicant claims small entity status. See 37 CFR 1.27

TOTAL AMOUNT OF PAYMENT (\$ 200)

## Complete if Known

Application Number	09/943,648
Filing Date	Aug 30, 2001
First Named Inventor	Dr. M. EL-Nagger
Examiner Name	B. Kwon
Group Art Unit	1614
Attorney Docket No.	

## METHOD OF PAYMENT (check all that apply)

Check  Credit card  Money  Other  None

Deposit Account: Visa / Exp. date 08/04

Deposit Account  
Number  
Deposit  
Account  
Name

6052  
Shaker Mousa/M. EL-Nagger

The Commissioner is authorized to: (check all that apply)

- Charge fee(s) indicated below  Credit any overpayments  
 Charge any additional fee(s) during the pendency of this application  
 Charge fee(s) indicated below, except for the filing fee  
 to the above-identified deposit account.

## FEE CALCULATION

## 1. BASIC FILING FEE

Large Entity	Small Entity	Fee Description	Fee Paid
Fee Code (\$)	Fee Code (\$)		
101 740	201 370	Utility filing fee	
106 330	206 165	Design filing fee	
107 510	207 255	- Plant filing fee	
108 740	208 370	Reissue filing fee	
114 160	214 80	Provisional filing fee	

SUBTOTAL (1) (\$ )

## 2. EXTRA CLAIM FEES FOR UTILITY AND REISSUE

Total Claims	-20** =	X	=
Independent Claims	-3** =	X	=
Multiple Dependent			

Large Entity	Small Entity	Fee Description
Fee Code (\$)	Fee Code (\$)	
103 18	203 9	Claims in excess of 20
102 84	202 42	Independent claims in excess of 3
104 280	204 140	Multiple dependent claim, if not paid
109 84	209 42	** Reissue independent claims over original patent
110 18	210 9	** Reissue claims in excess of 20 and over original patent

SUBTOTAL (2) (\$ )

\*\*or number previously paid, if greater; For Reissues, see above

## 3. ADDITIONAL FEES

Large Entity/Small Entity

Fee Code (\$)	Fee Code (\$)	Fee Description	Fee Paid
105 130	205 65	Surcharge - late filing fee or oath	
107 510	227 25	Surcharge - late provisional filing fee or cover sheet	
133 150	139 130	Non-English specification	
147 2,520	147 2,520	For filing a request for ex parte reexamination	
112 920*	112 920*	Requesting publication of SIR prior to Examiner action	
113 1,840*	113 1,840*	Requesting publication of SIR after Examiner action	
115 110	215 55	Extension for reply within first month	
116 400	216 200	Extension for reply within second month	200
117 920	217 460	Extension for reply within third month	
118 1,440	218 720	Extension for reply within fourth month	
128 1,960	228 980	Extension for reply within fifth month	
119 320	219 160	Notice of Appeal	
120 320	220 160	Filing a brief in support of an appeal	
121 280	221 140	Request for oral hearing	
138 1,510	138 1,510	Petition to institute a public use proceeding	
140 110	240 55	Petition to revive - unavoidable	
141 1,280	241 640	Petition to revive - unintentional	
142 1,280	242 640	Utility issue fee (or reissue)	
143 460	243 230	Design issue fee	
144 620	244 310	Plant issue fee	
122 130	122 130	Petitions to the Commissioner	
123 50	123 50	Processing fee under 37 CFR 1.17(q)	
126 180	126 180	Submission of Information Disclosure Stmt	
581 40	581 40	Recording each patent assignment per property (times number of properties)	
146 740	246 370	Filing a submission after final rejection (37 CFR § 1.129(a))	
149 740	249 370	For each additional invention to be examined (37 CFR § 1.129(b))	
179 740	279 370	Request for Continued Examination (RCE)	
169 900	169 900	Request for expedited examination of a design application	

Other fee (specify) \_\_\_\_\_

\*Reduced by Basic Filing Fee Paid

SUBTOTAL (3) (\$ )

200

## SUBMITTED BY

Name (Print/Type)	Registration No. (Attorney/Agent)	Telephone
Mawahel EL-Nagger		610-869-9358
Signature	Mawahel EL-Nagger	Date 09/10/02

Complete (if applicable)

WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.

Burden Hour Statement: This form is estimated to take 0.2 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, DC 20231.



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

RECEIVED  
OCT 03 2002

TECH CENTER 1600/2900

Application of : El-Naggar et al.

Examiner: B. Kwon

Appl. No. : 09/943,048

Art Unit: 1614

Filed : August 30, 2001

For : Treatment of Inflammatory, Cancer, and Thrombosis Disorders

Commissioner of Patents  
And Trademarks  
Washington, D.C. 20231

Sir:

This Amendment responds to the Office Action mailed April 10, 2002. A request for two months extension of time accompanies this Amendment. Please amend the application as follows:

**Response to Restriction Requirement**

Applicants affirm, with traverse, election of group III (Claim 4) in response to a restriction requirement.

**Amendment**

**IN THE CLAIMS:**

Please amend claims 1 and 5 as follows:

1. (Amended) A method of treating inflammatory disorders in a mammal comprising administering to said mammal a combination of: (i) a standard therapeutic dose of a COX2 inhibitor selected from the group consisting of [(i)] celecoxib [(CelebrexR) [A,]] and rofecoxib [(VioxxR) [B], and other specific COX2 inhibitors [C]], (ii) low dose aspirin in the amount of [(] 70-85 mg[), and (iii) antioxidant flavanoids, flavonoids or isoflavones.
  
5. (Amended) The method of claim 1, wherein the [COX2 inhibitors is compound [A], compound [B] or compound [C] and] low dose aspirin [of 70-85 mg] is in an enteric coated formulation. [This preferred enteric-coated formulation should allow for improved efficacy and safety.]